

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (currently amended): A method for analyzing the C-terminal amino acid sequence of a peptide to be examined, which method comprises the following steps:

a step of preparing a mixture containing a series of reaction products that are obtained from the peptide to be examined by releasing the C-terminal amino acids successively by chemical means,

a step of analyzing the differences in molecular weight between said series of reaction products and the original peptide by means of mass spectrometry to measure the decreases in molecular weight associated with the successive release of the C-terminal amino acids, and

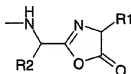
a step of identifying a series of the amino acids removed successively, based on a series of the measured decreases in molecular weight and arranging them from the C-terminus to obtain the information of the C-terminal amino acid sequence of the peptide,

wherein said process for releasing the C-terminal amino acids successively comprises at least the following steps:

a pretreatment step, for providing the protection by means of N-acylation, of allowing an alkanolic acid anhydride and an alkanolic acid both of vapor phase, which are supplied from a mixture of the alkanolic acid anhydride with a small amount of the alkanolic acid added thereto, to act on a dry sample of said peptide to be examined in a dry atmosphere at a temperature selected in a range of 10°C to 60°C and, thereby, applying, to the N-terminal amino group of the peptide

as well as to the amino group on the side chain of the lysine residue which may be included in the peptide, N-acylation by the acyl group derived from the alkanolic acid anhydride,

a step of allowing an alkanolic acid anhydride and a perfluoroalkanoic acid both of vapor phase, which are supplied from a mixture of the alkanolic acid anhydride with a small amount of the perfluoroalkanoic acid added thereto, to act on the dry peptide sample after N-acylation protection in a dry atmosphere at a temperature selected in a range of 15°C to 60°C and, thereby, releasing the C-terminal amino acids successively in association with a process that at the C-terminus of the peptide, the formation of a 5-oxazolone structure represented by the following general formula (III):



(III)

wherein R1 is a side chain of the C-terminal amino acid of the peptide and R2 is a side chain of the amino acid residue positioned just before the C-terminal amino acid, is followed by the cleavage of the 5-oxazolone ring, and

a hydrolysis treatment step which comprises applying, to a mixture containing a series of reaction products obtained in said step of releasing the C-terminal amino acids successively, a post-treatment of removing the remaining alkanolic acid anhydride and perfluoroalkanoic acid in a dry state, and then supplying vapor of a basic nitrogen-containing aromatic compound or a tertiary amine compound and water molecules, all of vapor phase, the vapor being generated from an aqueous solution dissolving the basic nitrogen-containing, aromatic compound or the tertiary amine compound therein, to allow the water molecules to act on the peptides of the

reaction products in the presence of the basic nitrogen-containing organic compound to give rise to the hydrolysis treatment, and after that conducting the re-dried up treatment by removing, from the mixture containing a series of reaction products, the remaining basic nitrogen-containing organic compound and water molecules to dry up the mixture,

wherein said step of measuring the decreases in molecular weight associated with the successive release of the C-terminal amino acids employs a technique which comprises:

allowing trypsin to act on said mixture, after the re-dried up treatment, containing a series of the reaction products finished by hydrolysis treatment, in a buffer solution, to carry out the treatment for the enzymatic digestion specific to trypsin of said peptide chain which holds N-acylation protection as for the N-terminal amino group of the peptide chain as well as to the amino group on the side chain of the lysine residue that may be contained in the peptide chain, and thereby, conducting selective cleavage of the C-terminal side peptide bond of each arginine residue that is present in the peptide chain to complete peptide fragmentization,

applying a desalting treatment to remove the buffer solution component, followed by recovering and drying the peptide fragments after the digestion treatment by trypsin, followed by drying,

next to that, conducting, as for the dried mixture containing said peptide fragments recovered after the digestion treatment by trypsin, molecular weight measurement for the cationic species of $(M+H)^+$ as well as molecular weight measurement for the anionic species of $(M-H)^-$, both of which are generated from the ionization treatment, by means of MALDI-TOF-MS,

with respect to the corresponding ion species, which are measured in said molecular weight measurement for the cationic species as well as molecular weight measurement for anionic species,

judging that the peaks of the peptide fragments each having an arginine residue at the C-terminus, which fragments are produced by said digestion treatment by trypsin, are peaks that give such intensities that the intensity in the molecular weight measurement for the cationic species of $(M+H)^+$ is relatively larger in comparison with the intensity in the molecular weight measurement for the anionic species of $(M-H)^-$, and judging that the peaks of the C-terminal peptide fragment derived from the original peptide and the C-terminal peptide fragments derived from a series of the reaction products that are obtained by successive release of the C-terminal amino acids, which fragments are produced by said digestion treatment by trypsin, are peaks that give such intensities that the intensity in the molecular weight measurement for the anionic species of $(M-H)^-$ is relatively larger in comparison with the intensity in the molecular weight measurement for the cationic species of $(M+H)^+$, and

based on a series of the peaks that gives a relatively larger intensity in the molecular weight measurement for the anionic species of $(M-H)^-$, measuring the decreases in molecular weight associated with the successive release of the C-terminal amino acids.

2. (original): A method claimed in Claim 1, wherein a symmetric anhydride of an alkanolic acid having 2 to 4 carbon atoms is used as the alkanolic acid anhydride contained in the mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto.

3. (original): A method claimed in Claim 2, wherein a symmetric anhydride of a linear chain alkanolic acid having 2 to 4 carbon atoms is used as the symmetric anhydride of said alkanolic acid having 2 to 4 carbon atoms.

4. (original): A method claimed in Claim 1, wherein acetic anhydride is used as the alkanolic acid anhydride contained in said mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto.

5. (original): A method claimed in Claim 1, wherein a perfluoroalkanoic acid of which a pKa is in the range of 0.3 to 2.5 is used as the perfluoroalkanoic acid contained in said mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto.

6. (original): A method claimed in Claim 1, wherein a perfluoroalkanoic acid having 2 to 4 carbon atoms is used as the perfluoroalkanoic acid contained in said mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto.

7. (original): A method claimed in Claim 6, wherein a linear chain perfluoroalkanoic acid having 2 to 4 carbon atoms is used as the perfluoroalkanoic acid having 2 to 4 carbon atoms.

8. (original): A method claimed in Claim 1, wherein a content of the perfluoroalkanoic acid in the mixture of an alkanolic acid anhydride with a small amount of a

perfluoroalkanoic acid added thereto is selected in a range of 1 to 20% by volume relative to the total volume of the alkanolic acid anhydride and the perfluoroalkanoic acid.

9. (Original): A method claimed in Claim 1, wherein, in the treatment using said mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto, said dry atmosphere is in a state that not only water but also oxygen have been eliminated.

10. (original): A method claimed in Claim 9, wherein said dry atmosphere is attained by, in an air-tight container, evacuating the inside atmosphere.

11. (original): A method claimed in Claim 1, wherein, in the treatment using said mixture of an alkanolic acid anhydride with a small amount of a perfluoroalkanoic acid added thereto, the temperature used is a temperature selected in a range of 15°C to 50°C.

12. (currently amended): A method for analyzing the C-terminal amino acid sequence of a peptide to be examined, which method comprises the following steps:

a step of preparing a mixture containing a series of reaction products that are obtained from the peptide to be examined by releasing the C-terminal amino acids successively by chemical,

a step of analyzing the differences in molecular weight between said series of reaction products and the original peptide by means of mass spectrometry to measure the decreases in molecular weight associated with the successive release of the C-terminal amino acids, and

a step of identifying a series of the amino acids removed successively, based on a series of the measured decreases in molecular weight and arranging them from the C-terminus to obtain the information of the C-terminal amino acid sequence of the peptide,

wherein said process for releasing the C-terminal amino acids successively, for the sample of the target peptide that has been subjected to separation by gel electrophoresis and is maintained in a state that it is bound on a polyacrylamide gel carrier, comprises the following steps:

a step of removing the water solvent impregnated into the polyacrylamide gel carrier by dilution with use of a polar aprotic solvent having no solvency for the polyacrylamide gel substance and having affinity for water, to conduct a dehydration treatment for the gel carrier,

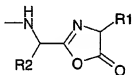
a pretreatment step for the target peptide sample that is still bound on the polyacrylamide gel carrier after carrying out said step for dehydration treatment, in which pretreatment step

applying N-acylation protection by the acyl group derived from the alkanolic acid constituting said alkanolic acid anhydride, to the N-terminal amino group of the target peptide with use of a solution of the alkanolic acid anhydride dissolved in a dipolar aprotic solvent that is capable of infiltrating into the polyacrylamide gel substance and keeping it in a swollen state is conducted by immersing, at a temperature selected in a range of 30°C to 80°C, the polyacrylamide gel carrier in the solution of the alkanolic acid anhydride to allow the alkanolic acid anhydride to act on the target peptide sample that is kept in the bound state; and then

removal of said solution is carried out by dilution with use of a polar aprotic solvent having no solvency for the polyacrylamide gel substance and having affinity for the alkanolic acid anhydride as well as the dipolar aprotic solvent, to conduct termination of the N-acylation reaction and removal of the reaction reagent therefor;

a step of treatment for the target peptide sample bound on the polyacrylamide gel carrier, after the pretreatment step of N-acylation protection, comprising steps of:

immersing, at a temperature selected in a range of 30 °C to 80 °C, the gel carrier in a mixed solution of an alkanolic acid anhydride added with a small amount of a perfluoroalkanoic acid in relative ratio thereto dissolved in a dipolar aprotic solvent that is capable of infiltrating into the polyacrylamide gel substance and keeping it in a swollen state, to allow the alkanolic acid anhydride and the perfluoroalkanoic acid to act on the target peptide sample being kept in the bound state; thereby, successive release of the C-terminal amino acids results from the reaction process with use of the mixed solution in which the formation of a 5-oxazolone-ring structure represented by the following general formula (III):



(III)

wherein R1 is a side chain of the C-terminal amino acid of the peptide and R2 is a side chain of the amino acid residue positioned just before the C-terminal amino acid, is followed by the cleavage of the 5-oxazolone-ring, and

removing the mixed solution used in the reaction for successive release of C-terminal amino acids, by dilution with use of a polar aprotic solvent having no solvency for polyacrylamide the gel substance and having affinity for the perfluoroalkanoic acid and the alkanolic acid anhydride as well as the dipolar aprotic solvent, to conduct termination of the releasing reaction and removal of the reaction reagents therefor; and

an additional step for hydrolysis treatment and then re-dehydration treatment, in which step

the hydrolysis treatment for said mixture comprising a series of reaction products obtained by the reaction for successive release of C-terminal amino acids is conducted by immersing the polyacrylamide gel carrier in an aqueous solution dissolving a basic nitrogen-containing aromatic compound or a tertiary amine compound therein to allow a water molecule to act, in the presence of said basic nitrogen-containing organic compound, on said peptides of the reaction products being still bound on the polyacrylamide gel carrier, and then,

the re-dehydration treatment for the gel carrier is performed by removing said aqueous solution infiltrated into the polyacrylamide gel carrier by dilution with use of a polar aprotic solvent having no solvency for the gel substance and having affinity for water; and

wherein said step of measuring the decreases in molecular weight associated with the successive release of the C-terminal amino acids employs a technique which comprises:

allowing trypsin to act on said mixture, after the re-dried up treatment, containing a series of the reaction products finished by hydrolysis treatment, in a buffer solution, to carry out the treatment for the enzymatic digestion specific to trypsin of said peptide chain which holds N-acylation protection as for the N-terminal amino group of the peptide chain as well as to the amino group on the side chain of the lysine residue that may be contained in the peptide chain, and thereby, conducting selective cleavage of the C-terminal side peptide bond of each arginine residue that is present in the peptide chain to complete peptide fragmentation,

applying a desalting treatment to remove the buffer solution component, followed by recovering and drying the peptide fragments after the digestion treatment by trypsin, followed by drying,

next to that, conducting, for the dried mixture containing said peptide fragments recovered after the digestion treatment by trypsin, molecular weight measurement for the

cationic species of $(M+H)^+$ as well as molecular weight measurement for the anionic species of $(M-H)^-$, both of which are generated from the ionization treatment, by means of MALDI-TOF-MS,

with respect to the corresponding ion species, which are measured in said molecular weight measurement for the cationic species of $(M+H)^+$ as well as molecular weight measurement for the anionic species of $(M-H)^-$,

judging that the peaks of the peptide fragments each having an arginine residue at the C-terminus, which fragments are produced by said digestion treatment by trypsin, are peaks that give such intensities that the intensity in the molecular weight measurement for the cationic species of $(M+H)^+$ is relatively larger in comparison with the intensity in the molecular weight measurement for the anionic species of $(M-H)^-$, and judging that the peaks of the C-terminal peptide fragment derived from the original peptide and the C-terminal peptide fragments derived from a series of the reaction products that are obtained by successive release of the C-terminal amino acids, which fragments are produced by said digestion treatment by trypsin, are peaks that give such intensities that the intensity in the molecular weight measurement for the anionic species of $(M-H)^-$ is relatively larger in comparison with the intensity in the molecular weight measurement for the cationic species of $(M+H)^+$, and

based on a series of the peaks that gives a relatively larger intensity in the molecular weight measurement for the anionic species of $(M-H)^-$, measuring the decreases in molecular weight associated with the successive release of the C-terminal amino acids.

13. (original): A method claimed in Claim 12, wherein a symmetric anhydride of an alkanolic acid having 2 to 4 carbon atoms is used as the alkanolic acid anhydride contained in the

mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanolic acid anhydride is dissolved.

14. (original): A method claimed in Claim 13, wherein a symmetric anhydride of a linear chain alkanolic acid having 2 to 4 carbon atoms is used as the symmetric anhydride of said alkanolic acid having 2 to 4 carbon atoms.

15. (original): A method claimed in Claim 12, wherein acetic anhydride is used as the alkanolic acid anhydride contained in the mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanolic acid anhydride is dissolved.

16. (original): A method claimed in Claim 12, wherein a perfluoroalkanoic acid of which a pKa is in the range of 0.3 to 2.5 is used as the perfluoroalkanoic acid contained in the mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanolic acid anhydride is dissolved.

17. (original): A method claimed in Claim 12, wherein a perfluoroalkanoic acid having 2 to 4 carbon atoms is used as the perfluoroalkanoic acid contained in the mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanolic acid anhydride is dissolved.

18. (original): A method claimed in Claim 17, wherein a linear chain perfluoroalkanoic acid having 2 to 4 carbon atoms is used as the perfluoroalkanoic acid having

2 to 4 carbon atoms in the mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanoic acid anhydride is dissolved.

19. (original): A method claimed in Claim 12, wherein, in the mixed solution where a small amount of the perfluoroalkanoic acid in relative ratio to the alkanoic acid anhydride is dissolved, the content ratio of the alkanoic acid anhydride and the perfluoroalkanoic acid is selected in the range of 1 to 20 volumes of the perfluoroalkanoic acid per 100 volumes of the alkanoic acid anhydride.